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Search Results -

Terms	Documents
115 same (advantag$ or useful$)	0

Database:

US Patents Full-Text Database
US Pre-Grant Publication Full-Text Database
JPO Abstracts Database
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Derwent World Patents Index
IBM Technical Disclosure Bulletins

Refine Search:

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115 same (advantag$ or useful$)
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| Clear |

Search History

Today's Date: 4/30/2001

DB Name	Query	Hit Count	Set Name
USPT	l15 same (advantag$ or useful$)	0	L16
USPT	l11 same translat$	3	L15
USPT	l11 same (advantag$ or useful$)	0	L14
USPT	l12 same (advantag$ or useful$)	0	L13
USPT	l11 same vector$	3	L12
USPT	capping near0 independent	9	L11
USPT	capping near0 independent near0 manner	0	L10
USPT	l8 same (advantag$ or useful$)	1	L9
USPT	l7 same (expression near0 vector)	11	L8
USPT	IRES near0 sequence	81	L7
USPT	l5 same (advantag$ or useful$)	15	L6
USPT	l4 same cleavage	607	L5
USPT	SpeI or SalI	5586	L4
USPT	TCR near0 expression near0 cassette	1	L3
USPT	TCR near0 beta near0 expression near0 cassette	0	L2
USPT	TCR near0 alpha near0 expression near0 cassette	0	L1

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Search Results -

Terms	Documents
l3 same (T near0 cell near0 receptor)	4

Database:

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US Patents Full-Text Database
US Pre-Grant Publication Full-Text Database
JPO Abstracts Database
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Refine Search:

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l3 same (T near0 cell near0 receptor)
```

| Clear |

Search History

Today's Date: 4/30/2001

DB Name	Query	Hit Count	Set Name
USPT	l3 same (T near0 cell near0 receptor)	4	L4
USPT	l2 same TCR	41	L3
USPT	expression near0 vector$	17168	L2
USPT	TCR near0 expression near0 vector$	0	L1

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Search Results -

Terms	Documents
l1 same (advantag$ or useful$)	10

Database:

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US Patents Full-Text Database
US Pre-Grant Publication Full-Text Database
JPO Abstracts Database
EPO Abstracts Database
Derwent World Patents Index
IBM Technical Disclosure Bulletins
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```
l1 same (advantag$ or useful$)
```

| Refine Search: | | Clear |

Search History

Today's Date: 4/30/2001

DB Name	Query	Hit Count	Set Name
USPT	l1 same (advantag$ or useful$)	10	L2
USPT	IRES near0 sequence$	84	L1

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☐ | Generate Collection

L4: Entry 3 of 4   File: USPT   Aug 17, 1999

DOCUMENT-IDENTIFIER: US 5939400 A
TITLE: DNA vaccination for induction of suppressive T cell response

DEPR:
The V.beta.8.2 gene of the T cell receptor was cloned in an expression vector
and injected three times at weekly intervals into the tibialis anterior muscle
of PL/J female mice, beginning one week following a single intramuscular
injection of cardiotoxin. Control PL/J mice were immunized once with
cardiotoxin and then at weekly intervals with the DNA coding for the
V.beta.5.1 variable region of the TCR. This TCR V region is not found on
pathogenic T cell clones in H-2.sup.u mice. Following this immunization
protocol, all mice were immunized with complete Freund's adjuvant containing
either the myelin basic protein peptide Ac1-20, or guinea pig MBP. The mice
vaccinated with the V.beta.8.2 DNA were resistant to experimental autoimmune
encephalitis (EAE) induced by the Ac1-20 peptide, while EAE was induced in the
groups vaccinated with the control V.beta.5.1 DNA, or with cardiotoxin alone.
Only one mouse out of 14 vaccinated with the V.beta.8.2 DNA developed mild
paralysis of the hind limbs, while 11/15 and 10/13 mice were paralyzed or died
in the respective control groups ($p < 0.0001$), following immunization with
pAc1-20. The data is shown in Table 1 and FIG. 1A.

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End of Result Set

☐   Generate Collection

L4: Entry 4 of 4   File: USPT   Aug 4, 1998

DOCUMENT-IDENTIFIER: US 5789208 A
TITLE: Polyclonal antibody libraries

DEPR:
Another embodiment of the invention is directed to a method for creating a
library of receptor proteins or any proteins which show variability. Receptor
proteins which may be utilized in this method may be any eukaryotic or
prokaryotic proteins which have variable regions including T-cell receptors
such as the TcR, B-cell receptors including immunoglobulins, natural killer
cell (NK) receptors, macrophage receptors and parts and combinations thereof.
Briefly, a sample of biological tissue, such as normal tissue, neoplastic
tissue, infected tissue, tissues containing extracellular matrix (ECM)
proteins, or any abnormal tissue, is introduced to a cell population capable
of producing the receptor proteins. The cell population is fixed and the cells
permeabilized. The variable region mRNAs of the receptor proteins are reverse
transcribed into cDNA sequences using a reverse transcriptase. The cDNA
sequences are PCR amplified and linked, preferably by hybridization of
complementary sequences at the terminal regions of these cDNAs. The linked
sequences are PCR amplified to create a population of DNA fragments which
encode the variable regions of the receptor proteins. These DNA fragments
contain the variable regions linked, preferably, in a head-to-head
transcriptional orientation and are cloned in-mass into expression vectors.
Useful expression vectors include phages such as display phages, cosmids,
viral vectors, phagemids or combinations thereof, and the vectors transformed
into host organisms and the different populations of organisms expanded. The
expression vectors which encode the recombinant receptor proteins are selected
and the sub-population expanded. The sub-population may be subcloned into
expression vectors, if necessary, which contain receptor constant region genes
in-frame and the library again expanded and expressed to produce the
sub-library of selected receptor proteins. Chimeric libraries can be easily
created by cloning the selected variable region genes into expression vectors
containing constant region genes of other proteins such as antibody constant
region genes or T cell receptor genes. The selected sub-libraries can be used
directly or transferred to other expression vectors before transfection into
host cells. Host cells may be T cells derived from the patient which, when
introduced back into the patient, express the receptor library on their
surface. This type of T cell therapy can be used to stimulate an immune
response to treat the same diseases as those described for antibody therapy.

```
=> s  TCR(w)expression (w) vector

L1              4 TCR(W) EXPRESSION (W) VECTOR

=> d bib ab l1 1-4
```

L1   ANSWER 1 OF 4   MEDLINE
AN   1999008315   MEDLINE
DN   99008315   PubMed ID: 9794213
TI   A retroviral vector that directs simultaneous expression of alpha and beta
     T cell receptor genes.
AU   Pogulis R J; Pease L R
CS   Department of Immunology, Mayo Foundation for Education and Research,
     Rochester, MN 55905, USA.
SO   HUMAN GENE THERAPY, (1998 Oct 10) 9 (15) 2299-304.
     Journal code: A12; 9008950. ISSN: 1043-0342.
CY   United States
DT   Journal; Article; (JOURNAL ARTICLE)
LA   English
FS   Priority Journals
EM   199901
ED   Entered STN: 19990115
     Last Updated on STN: 19990115
     Entered Medline: 19990105

AB   The transfer of alpha/beta T cell receptor (TCR) genes into T lymphocytes
     or their precursors could provide a means to increase frequency of tumor-
     or pathogen-specific cytotoxic T lymphocytes. To begin to address this
     possibility, we have used class I MHC-restricted alpha/beta TCR cDNAs to
     develop a retroviral **TCR expression vector**.
     Alpha- and beta-chain cDNAs were inserted into a derivative of the LN
     series of retroviral vectors, with the retroviral LTR directing expression
     of TCR-beta and an internal cytomegalovirus promoter/enhancer driving
     TCR-alpha expression. The variable region fragments can be replaced using
     unique restriction sites that have been introduced into the proximal
     constant regions. We have used this vector system to transfer two
     different pairs of alpha/beta TCR genes into an alpha- and
     beta-chain-deficient T cell hybridoma. TCR- hybridoma cells were
     transduced by coculture with pools of virus-producing cells, and
     fluorescence-activated cell sorting was used to enrich for cells
     expressing the transduced TCR. Transduction with either alpha/beta TCR
     restores stable, long-lived expression of the alpha/beta TCR complex.
     TCR-mediated signal transduction is also reconstituted, as demonstrated by
     the ability of transduced cells to secrete IL-2 following stimulation with
     Vbeta-specific antibodies. Our results suggest that alpha/beta T cell
     receptor gene transfer could provide a basis for new approaches to
     immunotherapy, and that further studies examining the in vivo fate of
     transduced TCR are possible.